Application No. 10/635,294 Applicant: Ilse Zolle

In the Claims:

- 1. (Cancelled)
- 2. (Currently Amended) The A compound of claim 1, having the formula (IA)

(IA)

wherein

R¹ is linear or branched C₁-C₄ alkyl, optionally substituted with a halogen selected from F, CL, I or Br;

R² denotes an alkyl group containing 1 or 2 carbon atoms; and

X is a halogen selected from the group consisting of I, BR, Cl and F a radioactive halogen selected from the group consisting of ¹²³I, ¹²⁴I, ¹²⁵I ¹³¹I, ⁷⁶Br, ⁸²Br or ¹⁸F.

- 3. (Cancelled)
- 4. (Currently Amended) The compound of claim ± 2 wherein R^1 and R^2 are each methyl, and X is non-radioactive or-radioactive iodine, and wherein the compound is I-metomidate (IMTO).

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- 5. (Currently Amended) The compound of claim $\frac{1}{2}$, wherein \mathbb{R}^1 is ethyl, \mathbb{R}^2 is methyl and X is non-radioactive or radioactive iodine, wherein the compound is I-iodometomidate (IMTO).
- 6. (Withdrawn) The compound of the formula (II)

wherein

- R¹ is linear or branched C₁-C₄ alkyl, optionally substituted with a halogen selected from the group consisting of F, Cl, I or Br;
- R² denotes an alkyl group containing 1 or 2 carbon atoms; and
- L represents an alkyl-stannyl group selected from the group consisting of a trimethylstannyl, triethylstannyl, tri-n-propylstannyl and tri-n-butylstannyl.
- 7. (Withdrawn) The compound of claim 6, having the general formula (IIA)

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wherein L is a trimethylstannyl group.

- 8. (Withdrawn) The compound of claim 6 wherein R¹ and R² are each methyl, and L is a trimethylstannyl group.
- 9. (Withdrawn) A process for preparing the compound of claim 1, the process comprising the steps of:
 - (a) providing a (S)-secondary alcohol of formula (III)

(b) coupling said (S)-secondary alcohol of formula (III) to an alkyl imidazole-5 [4]-carboxylate of formula (IV)

under conditions effective to achieve the compound of claim 1.

- 10. (Withdrawn) The process of claim 9, wherein the (S)-secondary alcohol of formula (III) is prepared by the process further comprising the steps of:
 - (a) reducing a substituted phenyl methyl ketone having X as either iodine or bromine, to racemic alcohol;
 - (b) preparing the chloroacetate of said racemic alcohol; and
 - (c) performing a lipase SAM II-catalysed resolution of (S)-alcohol of formula III derived from the (S)-enantiomeric ester.
- 11. (Withdrawn) A process for preparing the compound of claim 2, the process comprising the steps of
 - (a) preparing a compound of formula (II)
 - (b) reacting said compound of formula (II) under conditions effective for replacing L with non-radioactive or radioactive halogen to produce a compound of the formula (I) wherein R¹ is linear or branched C1-C4 alkyl, and is optionally substituted with a halogen selected from F, CL, I, Br; R² denotes an alkyl group containing 1 or 2 carbon atoms; and x is non-radioactive or radioactive halogen.
- 12. (Withdrawn) The compound of claim 4 having the structure ¹²³I-IMTO, ¹²³I-ETO, ¹²⁵I-IMTO, ¹²⁵I-ETO, ¹³¹I-IMTO, ¹³¹I-ETO, ¹²⁴I-IMTO, ¹²⁴I-ETO, ⁷⁶Br-MTO, ⁷⁶Br-ETO, ⁸²Br-ETO, ¹⁸F-MTO, ¹⁸F-ETO,, I-MTO (non-radioactive iodine), preferably ¹²³I-ETO or most preferably ¹³¹I-ETO.
- 13. (Withdrawn) The compound of claim 1, wherein X is a radioactive halogen, especially bromine.

- 14. (Currently amended) The compound of claim 1 2, wherein R1 is non-radioactive or radioactive 2-fluoroethyl, preferably radioactive.
- 15. (Withdrawn) A method for the in vivo detection of receptor positive tissue and tumors of adrenal cortex in persons with adrenal pathology, said method comprising administering the compound of claim 1 to said person with adrenal disease, and wherein a radiotracer is selected from the group consisting of gamma or positron-emitting halogens.
- 16. (Withdrawn) The method of claim 15, wherein the adrenal-derived turnor is not anatomically confined to the adrenal glands.
- 17. (Withdrawn) The compound of claim 5 having the structure ¹²³I-IMTO, ¹²³I-ETO, ¹²⁵I-IMTO, ¹²⁵I-ETO, ¹³¹I-IMTO, ¹³¹I-ETO, ¹²⁴I-IMTO, ¹²⁴I-ETO, ⁷⁶Br-MTO, ⁷⁶Br-ETO, ⁸²Br-ETO, I-MTO (non-radioactive iodine), ¹²³I-ETO or ¹³¹I-ETO.
- 18. (Canceled)